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TRANSPARENT AND HIGHLY HEAT-RESISTANT POLYIMIDE  
PRECURSOR AND PHOTORESISTIVE POLYIMIDE COMPOSITION  
THEREOF

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Technical Field

The present invention relates to an aqueous alkali developable photosensitive polyimide precursor resin 10 composition that is appropriate for highly heat-resistant transparent protection layers and insulation layers for liquid crystal display devices. In more detail, the present invention relates to a reactive transparent polyimide precursor represented in the following Chemical Formula 1, wherein it is 15 prepared by following steps of preparing transparent linear polyamic acid (A) from (a-1) one or more of tetracarboxylic acid dianhydrides selected from alicyclic tetracarboxylic acid dianhydrides having 3 to 30 carbon atoms, and (a-2) one or more of diamines selected from aliphatic, alicyclic, or non- 20 conjugated aromatic diamines of which number of carbon atoms is 3 to 30 and side chains have one or more ethylenically unsaturated bonds that may be crosslinked by a radical; and esterifying the transparent linear polyamic acid (A) thus obtained with an ethylenically unsaturated compound (B) containing 25 an epoxy group in the same molecule, and an aqueous alkali

developable negative-type photosensitive transparent polyimide precursor resin composition prepared by using the same as a main component.

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#### Background Art

Generally, polyimide resins are prepared by condensation polymerization of aromatic tetracarboxylic acid or its derivatives and aromatic diamine or aromatic diisocyanate, and polyimide resins thus obtained have superior heat resistance, chemical resistance, mechanical and electrical properties. Photosensitive aromatic polyimides having such superior properties have been widely used for electronic materials such as semiconductor encapsulant, etc.

15 However, aromatic polyimides are not appropriate for transparent protection layers or insulation layers for liquid crystal display devices in that they have low transmittance in the visible region, are colored in yellow or brown, and have a relatively high dielectric constant. Epoxy resins or acrylic resin compositions have been widely used as transparent protection layers or insulation layers for liquid crystal display devices. However, when they are processed at the temperature of 250°C or higher, severe discoloration and film contraction may occur and thus subsequent processing condition may be limited.

20 25 Therefore, it is preferable that they are used under the

AMENDED SHEET(ART. 34)

temperature of 230 °C or lower in view of their inherent heat resistance.

Accordingly, polyimide materials have been taken into consideration in order to meet heat resistance and transparency simultaneously, and some methods of preparation of polyimides by using aliphatic cyclic acid anhydrides to obtain transparent polyimide coating or film have been reported (Macromolecules, 5 1994. 27, 1117 and 1993, 26, 4961, Japanese Patent Laid-Open 1994. 27, 1117 and 1993, 26, 4961, Japanese Patent Laid-Open No. H9-95533, Japanese Patent Laid-Open No. 2001-330721).

10 However, in order to use such transparent polyimide resins as protection layers or insulation layers for liquid crystal display devices practically, the process of minute pattern formation by lithography process, which uses a separate photoresist, should be added after polyimide film formation on 15 substrate composed of glasses, etc.

While the use of photosensitive transparent polyimide resin compositions is advantageous in view of cost reduction since the above-described process can be simplified, the photosensitive transparent polyimide resin composition using 20 trans-1,4-diaminocyclohexane reported in Japanese Patent Laid-Open No. 2002-161136 is practically inapplicable due to 25 too slow photosensitivity.

The inventors of the present invention have reported photosensitive transparent polyimide resin composition in 25 Korean Patent Application No. 10-2002-074070, leaving

development margin and photosensitivity to be improved further.

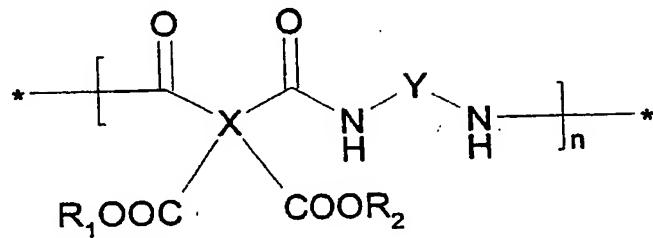
#### Disclosure of the Invention

5 In order to resolve the above-described problems, an object of the present invention is to provide a negative-type photosensitive transparent polyimide precursor having enough photosensitivity and a method for preparing the same, and to provide compositions that may be used for transparent protection 10 layers and insulation layers for liquid crystal display devices having superior heat resistance, chemical resistance, 15 mechanical strength, and electrical insulation.

The foregoing and other objects, aspects, and advantages would be better understood from the following detailed 15 description of preferred embodiments of the invention:

In order to achieve the above objects, the present invention provides a reactive transparent polyimide precursor represented by the following Chemical Formula 1:

20 [Chemical Formula 1]



wherein X is a tetra-valent organic group derived from

alicyclic tetracarboxylic acid dianhydrides having 3 to 30 carbon atoms;

Y is a di-valent organic group derived from aliphatic, alicyclic, or non-conjugated aromatic diamines of which number 5 of carbon atoms is 3 to 30 and side chains have one or more ethylenically unsaturated bonds that may be crosslinked by a radical; and

R<sub>1</sub> and R<sub>2</sub> are independently each other hydrogen atom, or organic group having 1 to 20 carbon atoms including one or more 10 ethylenically unsaturated bonds, provided that they are not hydrogen atom at the same time.

The acid value of the above reactive transparent polyimide precursors may be in the range of 30 to 200 mg KOH/g, and the molecular weight of the above reactive transparent polyimide 15 precursors may be in the range of 2,000 to 200,000.

Furthermore, the present invention provides a method for preparing reactive transparent polyimide precursors of the above Chemical Formula 1, comprising the steps of: preparing transparent linear polyamic acid (A) from (a-1) one or more of 20 tetracarboxylic acid dianhydrides selected from alicyclic tetracarboxylic acid dianhydrides having 3 to 30 carbon atoms, and (a-2) one or more of diamines selected from aliphatic, alicyclic, or non-conjugated aromatic diamines of which number 25 of carbon atoms is 3 to 30 and side chains have one or more ethylenically unsaturated bonds that may be crosslinked by a

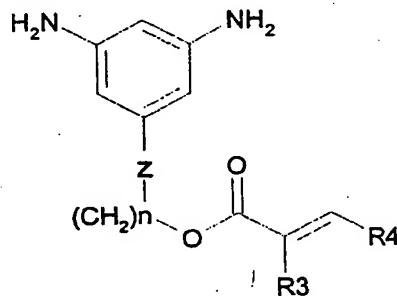
radical; and esterifying the transparent linear polyamic acid (A) thus obtained with an ethylenically unsaturated compound (B) containing an epoxy group in the same molecule.

The above tetracarboxylic acid dianhydrides may be one

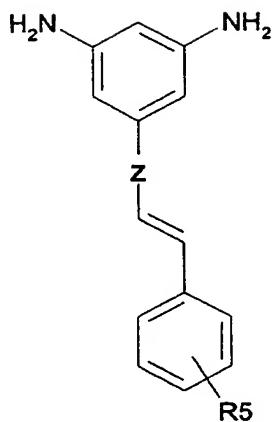
5 or more anhydride(s) selected from 1,2,3,4-cyclobutanetetracarboxylic acid dianhydride (CBDA), 1,2,3,4-cyclopentanetetracarboxylic acid dianhydride (CPDA), bicyclooctene-2,3,5,6-tetracarboxylic acid dianhydride (BODA), 5-(2,5-dioxotetrahydrofuran-3-yl)-3-methylcyclohexene-1,2-d 10 icarboxylic acid anhydride (DOCDA), 4-(2,5-dioxotetrahydrofuran-3-yl)-tetralin-1,2-dicarboxylic acid anhydride (DOTDA); and the above diamine may be one or more selected from those having the general chemical formulae represented by the following Chemical Formulae 7 to 9:

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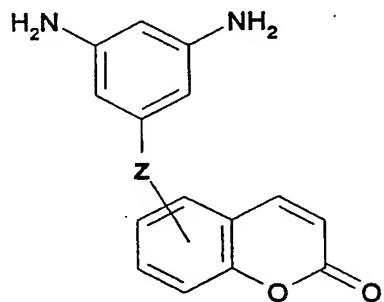
<Chemical Formula 7>



<Chemical Formula 8>



<Chemical Formula 9>



5 wherein Z is one of ester, amide, imide, ether, and carbonyl group; R<sub>3</sub>, R<sub>4</sub>, and R<sub>5</sub> are independently each other hydrogen atom or alkyl or aryl groups having 1 to 20 carbon atom(s); and n is an integer of 1 to 20.

10 The diamines in the above may be one or more selected from the group consisting of 2-(methacryloyloxy)ethyl 3,5-diaminobenzoate, 3,5-diaminophenyl cinnamate, and coumaronyl 3,5-diaminobenzoate.

The above ethylenically unsaturated compound (B) containing an epoxy group in the same molecule may be one or

more compounds selected from the group consisting of allyl glycidyl ether, glycidyl acrylate, glycidyl methacrylate, 3,4-epoxycyclohexylmethyl acrylate, 3,4-epoxycyclohexylmethyl methacrylate, glycidyl 5-norbornene-2-carboxylate (a mixture of endo and exo forms), glycidyl 5-norbornene-2-methyl-2-carboxylate (a mixture of endo and exo forms), 1,2-epoxy-5-hexene, and 1,2-epoxy-9-decene.

The present invention also provides photosensitive polyimide precursor resin compositions wherein they are prepared by using the above-mentioned transparent polyimide precursors and one or more photo-initiators as essential components, and adding one or more compounds selected from the group consisting of photosensitizers, multi-functional monomers, and common coating additives if necessary.

The portion of the above-described reactive transparent polyimide precursors may be 10 to 99 weight %, and that of the photo-initiators may be 0.1 to 90 weight %, based on the amount of total solids.

Thickness of polyimide films obtained by coating and postbaking the above composition may be 0.5 to 100  $\mu\text{m}$ ; thermal decomposition temperature of the polyimide films is in the range of 300 to 500°C; transmittance of the polyimide films 90% or higher between 400 to 700 nm; and dielectric constant of the polyimide films in the range of 2.5 to 4.0 measured at 1 kHz.

The present invention can also provide transparent protection layer or insulation layer, wherein it is prepared by using the photosensitive polyimide precursor resin compositions obtained in the above.

5 Further, the present invention can provide a liquid crystal display device wherein it is manufactured by applying the photosensitive polyimide precursor resin compositions obtained in the above to its transparent protection layer or insulation layer.

10 Still further, the present invention can provide a liquid crystal display device wherein it is manufactured by applying the photosensitive polyimide precursor resin compositions as the organic insulation materials for liquid crystal display devices.

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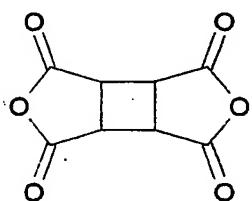
Hereinafter, the present invention is described in more detail as follows:

(a-1) is one or more of tetracarboxylic acid dianhydrides selected from alicyclic tetracarboxylic acid dianhydrides 20 having 3 to 30 carbon atoms, of which particular examples include, but are not limited to, 1,2,3,4-cyclobutanetetracarboxylic acid dianhydride (CBDA shown in Chemical Formula 2), 1,2,3,4-cyclopantanetetracarboxylic acid dianhydride (CPDA shown in Chemical Formula 3), 25 bicyclooct-7-ene-2,3,5,6-tetracarboxylic acid dianhydride

(BODA shown in Chemical Formula 4),  
 5-(2,5-dioxotetrahydrofuran-3-yl)-3-methylcyclohexene-1,2-dicarboxylic acid anhydride (DOCDA shown in Chemical Formula 5),  
 4-(2,5-dioxotetrahydrofuran-3-yl)-tetralin-1,2-dicarboxylic acid anhydride (DOTDA shown in Chemical Formula 6), etc.

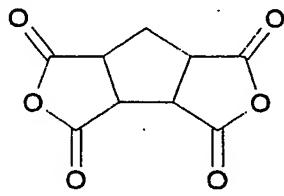
Also, if it is necessary, aromatic tetracarboxylic acid dianhydrides selected from the group consisting of pyromellitic acid dianhydride, benzophenone tetracarboxylic acid dianhydride, oxydiphtalic acid dianhydride,  
 10 hexafluoroisopropylidenediphtalic acid dianhydride, etc. may be further used as long as they do not deteriorate the transparency of the polyimides.

## [Chemical Formula 2]

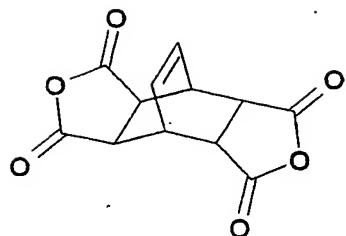


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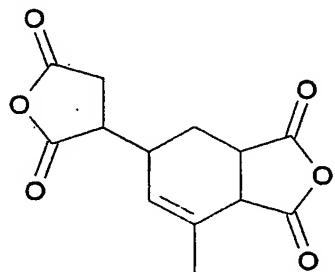
## [Chemical Formula 3]



20 [Chemical Formula 4]

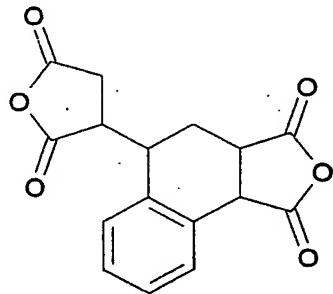


【Chemical Formula 5】



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【Chemical Formula 6】



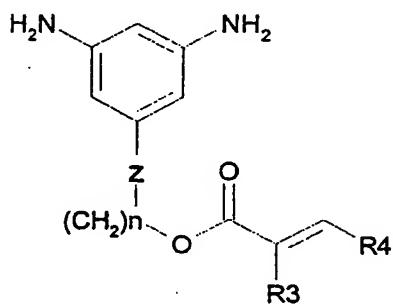
(a-2) is one or more of diamines selected from aliphatic, alicyclic, or non-conjugated aromatic diamines of which number of carbon atoms is 3 to 30 and side chains have one or more ethylenically unsaturated bonds that may be crosslinked by a radical, of which particular examples include 2-(methacryloyloxy)ethyl 3,5-diaminobenzoate (Chemical

Formula 7,  $R_3 = CH_3$ ,  $R_4 = H$ ,  $Z = COO$ ,  $n = 2$ ), 3,5-diaminophenyl cinnamate (Chemical Formula 8,  $R_5 = H$ ,  $Z = OCO$ ), coumaronyl 3,5-diaminobenzoate (Chemical Formula 9,  $Z = COO$ ), etc. (wherein  $Z$  is one of ester, amide, imide, ether, carbonyl group, etc.; 5  $R_3$ ,  $R_4$ , and  $R_5$  are independently each other hydrogen atom or alkyl or aryl group having 1 to 20 carbon atoms; and  $n$  is an integer of 1 to 20). Also, if necessary, diamines not including ethylenically unsaturated bonds at their side chains may be further used as long as they do not deteriorate the transparency 10 and chemical resistance of the polyimides thus formed.

Their particular examples include, but are not limited to, one or more compounds selected from the group consisting of 1,2-diaminocyclohexane, 1,3-diaminocyclohexane, 1,4-diaminocyclohexane, para-phenylenediamine (p-PDA), 15 meta-phenylenediamine (m-PDA), 4,4-oxydianiline (ODA), 4,4-methylenedianiline (MDA), 2,2-bis(aminophenyl) hexafluoropropane (HFDA), meta-bis(aminophenoxydiphenyl)sulfone (m-BAPS), para-bis(aminophenoxydiphenyl)sulfone (p-BAPS), 20 1,4-bis(aminophenoxy)benzene (TPE-Q), 1,3-bis(aminophenoxy)benzene (TPE-R), 2,2-bis(aminophenoxyphenyl)propane (BAPP), 2,2-bis(aminophenoxyphenyl)hexafluoropropane (HFBAPP), 5-diaminobenzoic acid, 2,4-diaminobenzenesulfonic acid, 25 2,5-diaminobenzenesulfonic acid,

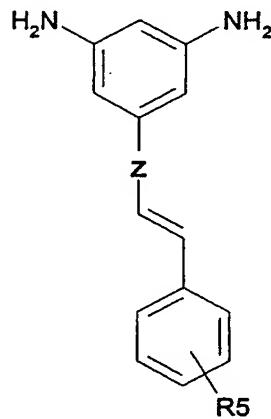
bis(aminopropyl)tetramethyldisiloxane, etc.

【Chemical Formula 7】

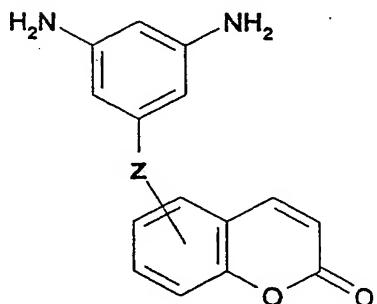


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【Chemical Formula 8】



【Chemical Formula 9】



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Hereinafter, a method for preparing a reactive transparent polyimide precursor according to the present invention is described as follows:

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<Step 1: Synthesis of transparent linear polyamic acid

(A) >

In the first step, the transparent linear polyamic acid (A) is prepared from tetracarboxylic acid dianhydrides of (a-1) 10 and diamines of (a-2) according to the conventional preparation method of polyamic acid. In other words, it is possible to obtain linear polyamic acid (A) quantitatively by adding tetracarboxylic acid dianhydride to diamine slowly, which is then reacted for 24 hours in a reactor, of which temperature 15 can be controlled, while maintaining the temperature below room temperature.

The solvent used for polymerization is one or more solvents selected from the group consisting of meta-cresol, N-methyl-2-pyrrolidone (NMP), dimethylformamide (DMF), 20 dimethylacetamide (DMAc),  $\gamma$ -butyrolactone, 2-butoxyethanol, and 2-ethoxyethanol. Organic acid such as para-toluene sulfonic acid, hydroxybenzoic acid, crotonic acid, etc. or an organic amine derivative, may be added in an amount of up to 5 weight % as an imidization catalyst relative to the total amount of the 25 above reaction mixture. Further, hydroquinone or similar

compound may be used as a thermopolymerization inhibitor in the range of 30 to 500 ppm relative to the weight of the polymer. Still further, adhesion may be improved by adding a siloxane diamine derivative within a range of 0.5 to 50 weight % with respect to the total amount of diamine, if necessary.

The average molecular weight (Mw) of the linear polyamic acid thus obtained is characterized by being in the range of 2,000 to 200,000. If the molecular weight is less than 2,000, it is difficult to form films thereafter; if it is higher than 10 200,000, it may be difficult to handle them due to excessively high viscosity, and the degree of planarization of the films formed thereafter may be worsened. The linear polyamic acid thus obtained shows photosensitivity when coupled with an appropriate photo-initiator without multi-functional monomer 15 since it has ethylenically unsaturated bonds at its polymer side chain. Also, it is advantageous to obtain films having superior chemical resistance after postbaking polyamic acid to obtain polyimide. However, the linear polyamic acid without subsequent treatment has a problem of narrow development margin 20 since its solubility in an alkali developer is very good.

<Step 2: Esterification with ethylenically unsaturated compound (B)>

In the second step, to control the solubility of the linear 25 polyamic acid in the alkali developer and to endow additional

reactive site, the alkali-soluble linear polyamic acid (A) is esterified with an ethylenically unsaturated compound (B) having an epoxy group in the same molecule. The compound (B) includes allyl glycidyl ether, glycidyl (meth)acrylate, 5 3,4-epoxycyclohexylmethyl (meth)acrylate, glycidyl 5-norbornene-2-carboxylate (a mixture of endo and exo forms), glycidyl 5-norbornene-2-methyl-2-carboxylate (a mixture of endo and exo forms), 1,2-epoxy-5-hexene, 1,2-epoxy-9-decene, etc., which may be used singly or in combination of two or more 10 compounds during esterification. It is preferable to use 10 to 400% equivalent weight of the compound (B) with respect to that of the total carboxylic acid group(s) in the linear polyamic acid (A). If this range is deviated, there may be problems such as difficulty in controlling the solubility to the developer 15 or reduced heat resistance of the films thus formed.

The acid value of the reactive transparent polyimide precursor thus obtained is about 30 to 200 mg KOH/g, and the average molecular weight is within a range of 2,000 to 200,000, and more preferably, within a range of 5,000 to 100,000.

20 The reactive transparent polyimide precursor obtained in the above as shown in Chemical Formula 1 has more ethylenically unsaturated bonds per structural unit compared to those of the linear polyamic acid (A), and therefore, it is possible to prepare negative-type photosensitive resin compositions with enough 25 development margin without additional multi-functional

monomers; to reduce the amount of photo-initiator; and to improve photosensitivity.

Hereinafter, a method for preparing negative-type photosensitive transparent polyimide precursor resin compositions with alkali developability by using the reactive transparent polyimide precursors obtained in the above is described.

In order to prepare a negative-type photosensitive transparent polyimide precursor resin composition with alkali developability, an appropriate photo-initiator (C) should be used, and if necessary, photosensitizer (D), multi-functional monomer (E), other additive (F), etc. may be used in combination with the above.

The above alkali-soluble reactive transparent polyimide precursor (shown in Chemical Formula 1) may be used singly or in combination with two or more. It is preferable to use the reactive transparent polyimide precursor in an amount of 10% to 99% of the total solid portion of the photosensitive transparent polyimide precursor resin compositions. If it is less than 10%, the adhesion of the films thus formed is decreased; if it is greater than 99%, there may be a problem of low photosensitivity.

Examples of the photo-initiator (C) include benzophenone, 4,4'-bis(N,N-diethylamino)benzophenone, 25 4,4'-bis(N,N-dimethylamino)benzophenone, phenylbiphenyl

ketone, 1-hydroxy-1-benzoylcyclohexane, benzil,  
 benzyldimethyl ketal,  
 2-benzyl-2-(dimethylamino)-4'-morpholinobutyrophenone,  
 2-methyl-4'-(methylthio)-2-morpholinopropiophenone,  
 5 thioxanthone, 1-chloro-4-propoxythioxanthone, isopropyl  
 thioxanthone, diethyl thioxanthone, ethyl anthraquinone,  
 4-benzoyl-4'-methyldiphenyl sulfide, benzoyl butyl ether,  
 2-hydroxy-2-benzoyl propane,  
 2-hydroxy-2-(4'-isopropyl)benzoyl propane,  
 10 4-butylbenzoyltrichloromethane, 4-phenoxy  
 benzoyle dichloromethane, benzoyl formic acid methyl,  
 1,7-bis(9'-acrydanyl)heptane,  
 9-n-butyl-3,6-bis(2'-morpholinoisobutyroyl)carbazole,  
 2,4,6-trimethylbenzoyl diphenylphosphin oxide, etc.  
 15 It is preferable to use 0.1 to 90 weight % of photo-initiator  
 relative to the amount of a polymer, and more preferably, within  
 a range of 0.2 to 50 weight %. If it deviates from this range,  
 there may be problems like no photo-polymerization, or reduced  
 developability, etc.  
 20 As a photosensitizer (D), a coumarine derivative such as  
 3,3'-carbonylbis(dimethylamino)coumarine, etc., an aromatic  
 carbonyl compound such as fluorenone, etc., an amine derivative  
 such as triethylamine, etc., a methylenethiazole derivative such  
 as 1,2-dihydronaphtho[1,2,d]thiazole, etc. may be used.  
 25 A multi-functional monomer (E) may be one of compounds

having one or more ethylenically unsaturated groups that may be additionally polymerized in a molecule and with a boiling point of 100°C or higher. Examples include phenoxyethyl methacrylate, polyethyleneglycol (meth)acrylate, 5 polypropyleneglycol (meth)acrylate, trimethylolethane tri(meth)acrylate, trimethylolpropane tri(meth)acrylate, neopentyl glycol (meth)acrylate, pentaerythritol triacrylate, pentaerythritol tetraacrylate, dipentaerythritol pentaacrylate, dipentaerythritol hexaacrylate, etc.

10 Also, functional monomers include KAYARAD DPCA-20, KAYARAD DPCA-30, KAYARAD DPCA-60, KAYARAD DPCA-120, etc. in the form of caprolactone-modified dipentaerythritol; KAYARAD TC-110S in the form of caprolactone-modified tetrahydrofurfuryl acrylate, KAYARAD HX-220, KAYARAD HK-620, etc. in the form of 15 caprolactone-modified neopentyl glycol hydroxypivalate, etc.

Other functional monomers include epoxy acrylates, novolak-epoxy acrylates that are bisphenol A derivatives; U-324A, U15HA, U-4HA, etc. that are urethane-based multi-functional acrylates. Functional monomers having ethylenically 20 unsaturated double bonds may be used singly or in combination with two or more monomers.

If necessary, other additives (F) that are generally used in coating agents can be included. Examples of F are thermopolymerization inhibitors such as hydroquinone, 25 4-methoxyphenol, quinone, pyrocatechol, t-butyl catechol,

phenothiazine, etc.; plasticizers; silicone-based adhesion promoters; fillers; surfactants; etc.

Described below is a method of pattern formation by using a negative-type photosensitive transparent polyimide precursor 5 resin composition of the present invention, and then forming a heat-resistant transparent polyimide film:

<Step 1>

A photosensitive polyimide precursor resin composition 10 is coated on substrate. For substrate, glass or transparent plastic resin can be used in view of the characteristics of liquid crystal display devices, but not limited as long as they meet the object of display. As to the method of coating the photosensitive resin composition of the present invention on 15 the surface of a substrate, various kinds of coating methods such as spraying, roll coating, slit nozzle coating, rotatory coating, extrusion coating, bar coating, etc. and also combination of two or more kinds of the above methods may be adopted. The thickness of coated films varies according to the 20 method of coating, concentration of solids in the composition, viscosity, etc., but is usually 0.5 to 100  $\mu\text{m}$  after they are dried.

<Step 2>

25 In order to form non-fluxionary film after coating, prebake

process, in which the solvent is volatilized under vacuum, infrared and/or heat, is performed. Conditions for heating vary according to the kind or mixing ratio of each component, but usually are in the range of 60 to 130°C for 5 to 500 seconds 5 on a hot plate, or 60 to 140°C for 20 to 1,000 seconds in an oven.

<Step 3>

Thereafter, the thin film thus obtained in step 2 is exposed 10 to radiation using a mask having a predetermined pattern. Exposure can be done by excimer laser, UV, visible light, e-beam, x-ray, etc. In the present invention, it is preferable to use g-line (having a wavelength of 436 nm), h-line (having a wavelength of 405 nm), and i-line (having a wavelength of 365 nm) of mercury lamp. Exposure can be carried out by contact, 15 proximity, or projection method.

<Step 4>

Whether to perform post-exposure bake (PEB) prior to alkali 20 development is optional. PEB step may be introduced if necessary. PEB temperature is normally 150°C or lower, and PEB time is about 0.1 to 10 minutes.

<Step 5>

25 In the development process, non-exposed part of the film

is removed by alkali developer. For the developer, an aqueous alkali solution consisting of mineral alkalis such as sodium hydroxide, potassium hydroxide, sodium carbonate, sodium silicate, sodium meta-silicate, ammonia, water, etc.; primary 5 amines such as ethylamine, n-propylamine, etc.; secondary amines such as diethylamine, di-n-propylamine, etc.; tertiary amines such as triethylamine, methyldiethylamine, n-methylpyrrolidone, etc.; alcohol amines such as dimethylethylalcohol amine, triethylalcohol amine, etc.; quarternary ammonium salts such 10 as tetramethyl ammonium hydroxide, tetraethyl ammonium hydroxide, choline, etc.; and amines such as pyrrole, piperidine, etc. may be used.

It is also possible to use an aqueous solution in which an appropriate amount of water-soluble organic solvent such as 15 methyl alcohol, ethyl alcohol, propyl alcohol, isopropyl alcohol, propylene glycol monomethyl ether, dipropylene glycol monomethyl ether, etc.; and surfactant; etc. is added to the above aqueous alkali solution. Development time is usually about 10 to 200 seconds. Dipping, spraying, puddle, etc. are applicable 20 for the development method. After development, deionized (DI) water rinse is done for 20 to 200 seconds, and moisture on the substrate is removed by spin drying or blowing with compressed air or nitrogen to give a desired pattern on the substrate.

After development, polyimide precursor films are converted into heat-resistant polyimide films through postbake (also called hardbake) on a hot plate or in an oven. Baking is usually done at 150°C to 300°C for 1 to 120 minutes on a hot plate, or 5 for 10 to 120 minutes at the same temperature range in an oven. Completely cross-linked and hardened polyimide pattern is obtained after postbake.

The thermal decomposition temperature of polyimide films thus formed is in the range of 300 to 500°C, and transmittance 10 between 400 to 700 nm is higher than 90%. It is also seen that the dielectric constant measured at 1 kHz is within a range of 2.5 to 4.0.

Heat-resistant polyimide films formed from the negative-type photosensitive polyimide precursor resin 15 compositions of the present invention can be used for protection layers, insulation layers, etc. of liquid crystal display devices.

Additional features and advantages of the invention will be set forth in the description which follows, and in part will 20 become apparent from the description, or may be learned by practice of the invention.

#### Modes for Carrying Out the Invention

25 First, polyimide precursors used in preferred embodiments

and comparative examples are synthesized in the following synthetic examples, and the linear polyamic acid used in the following synthetic examples is synthesized according to known method. This is also described in detail in Korean Patent 5 Application No. 10-2002-074070 which is invented by the present inventors. In other words, diamine is dissolved in solvent under nitrogen gas flow in a reactor to which a stirrer, temperature controller, nitrogen inlet, and cooler are attached, and stirred during slow addition of tetracarboxylic acid dianhydride, where 10 the reaction temperature is maintained below room temperature. The yield of the polymerization reaction is quantitative when the mixture is reacted for 12 ~ 24 hours.

<Synthetic Example 1>

15 A 100 g portion of the linear polyamic acid solution (solid content 20%, acid value of 220 mg KOH/g, Mw = 18,000) synthesized by reacting 1,2,3,4-cyclobutanetetracarboxylic acid dianhydride (CBDA) and 2-(methacryloyloxy)ethyl 3,5-diaminobenzoate (DA-HEMA) in N-methyl-2-pyrrolidinone 20 (NMP) /  $\gamma$ -butyrolactone (volume ratio = 1/1) co-solvent is charged in a 250 ml roundbottomed flask, and heated to 110°C under nitrogen atmosphere during stirring.

Reactive polyimide precursor 1 is synthesized by adding 20 g of glycidyl methacrylate to the flask slowly in an hour, 25 and allowing to react for two hours. The acid value of the

reactive polyimide precursor thus obtained is 120 mg KOH/g.

<Synthetic Example 2>

Reactive polyimide precursor 2 is synthesized in the same  
5 method as that in Synthetic Example 1 except that the linear  
polyamic acid (solid content 20%, acid value of 240 mg KOH/g,  
Mw = 22,000) is synthesized from 3,5-diaminophenyl cinnamate  
instead of 2-(methacryloyloxy)ethyl 3,5-diaminobenzoate in the  
above Synthetic Example 1. The acid value of the reactive  
10 polyimide precursor thus obtained is 160 mg KOH/g.

<Synthetic Example 3>

Reactive polyimide precursor 3 is synthesized in the same  
method as that in Synthetic Example 1 except that the linear  
15 polyamic acid (solid content 20%, acid value of 240 mg KOH/g,  
Mw = 22,000) is synthesized from coumaronyl 3,5-diaminobenzoate  
instead of 2-(methacryloyloxy)ethyl 3,5-diaminobenzoate in the  
above Synthetic Example 1. The acid value of the polyimide  
precursor thus obtained is 130 mg KOH/g.

20

<Synthetic Example 4>

Reactive polyimide precursor 4 is synthesized in the same  
method as that in Synthetic Example 1 except that the linear  
polyamic acid (solid content 20%, acid value of 200 mg KOH/g,  
25 Mw = 26,000) is synthesized from 4,4-oxydianiline (ODA) instead

of 2-(methacryloyloxy)ethyl 3,5-diaminobenzoate in the above Synthetic Example 1. The acid value of the polyimide precursor thus obtained is 140 mg KOH/g.

5                   <Synthetic Example 5>

Reactive polyimide precursor 5 is synthesized in the same method as that in Synthetic Example 1 except that the linear polyamic acid (solid content 20%, acid value of 230 mg KOH/g, Mw = 28,000) is synthesized from pyromellitic acid dianhydride (10 PMDA) and 4,4-oxydianiline instead of 1,2,3,4-cyclobutanetetracarboxylic acid dianhydride (CBDA) and 2-(methacryloyloxy)ethyl 3,5-diaminobenzoate in the above Synthetic Example 1. The acid value of the reactive polyimide precursor thus obtained is 140 mg KOH/g.

15

By using reactive polyimide precursors prepared in the above synthetic examples, photosensitive resin compositions in the following comparative examples and preferred embodiments are prepared:

20

[Preferred Embodiment 1]

Photosensitive polyimide precursor resin composition 1 is prepared by dissolving 0.5 g of CGI124 as a photo-polymerization initiator and 0.01 g of BYK307 as a 25 surfactant in 50 g of the reactive polyimide precursor solution

1 prepared in Synthetic Example 1 and filtering the mixture by a 0.2- $\mu\text{m}$  filter. Film is formed by spin-coating the above composition 1 on a glass substrate and pre-baking on a 100°C-hot plate for 90 seconds. A photomask is put onto the film thus obtained, and 20 mW/cm<sup>2</sup> UV light is irradiated at 365 nm for 10 seconds by using a mercury lamp. A pattern is formed by developing the film with 2.38 weight % tetramethylammonium hydroxide aqueous solution at room temperature for 20 seconds, and washing it with DI water for 10 seconds. It is then hardened by postbake step in an oven at 250°C for 1 hour. The minimum line width of the film thus obtained is 25  $\mu\text{m}$  (for 1:1 line/space pattern), and the film thickness is 1.2  $\mu\text{m}$ .

15 [Preferred Embodiment 2]

The same processes as those in Preferred Embodiment 1 are performed except that the reactive polyimide precursor solution 2 prepared in Synthetic Example 2 is used. The minimum line width of the film thus obtained is 25  $\mu\text{m}$ , and the film thickness is 1.2  $\mu\text{m}$ .

20

[Preferred Embodiment 3]

The same processes as those in Preferred Embodiment 1 are performed except that the reactive polyimide precursor solution 3 prepared in Synthetic Example 3 is used. The minimum line width of the film thus obtained is 30  $\mu\text{m}$ , and the film thickness

is 1.2  $\mu\text{m}$ .

[Comparative Example 1]

The same processes as those in Preferred Embodiment 1 are  
5 performed except that the reactive polyimide precursor solution  
4 prepared in Synthetic Example 4 is used. The minimum line  
width of the film thus obtained is 50  $\mu\text{m}$ , and the film thickness  
is 1.2  $\mu\text{m}$ .

10 [Comparative Example 2]

The same processes as those in Preferred Embodiment 1 are  
performed except that the reactive polyimide precursor solution  
5 prepared in Synthetic Example 5 is used. The minimum line  
width of the film thus obtained is 50  $\mu\text{m}$ , and the film thickness  
15 is 1.2  $\mu\text{m}$ .

[Comparative Example 3]

The same processes as those in Preferred embodiment 1 are  
performed by using the linear polyamic acid (solid content 20%,  
20 acid value of 220 mg KOH/g, Mw = 18,000) which is not reacted  
with glycidyl methacrylate in Synthetic Example 1 instead of  
the reactive polyimide precursor solution 1, where it is not  
possible to obtain a desired pattern having an appropriate shape,  
since the pattern is swept away in the process of development.  
25 In order to prevent pattern loss, post-exposure bake step at

160°C for 10 minutes should be added before development. Then, it becomes possible to form a desired pattern, where the minimum line width of the pattern thus obtained is 40  $\mu\text{m}$ , and the film thickness is 1.2  $\mu\text{m}$ .

5

[Testing Example]

Physical properties of polyimide films obtained from the photosensitive polyimide compositions in the above preferred embodiments are measured in a method described below, and the 10 results are shown in Table 1:

<Evaluation of heat resistance>

After the polyimide pattern is formed, the change in its thickness by thermal impact is monitored. It may be said that, 15 the smaller the change in its thickness at high temperature, the better the heat resistance of the film is. The ratio of the post-baked film thickness to pre-baked film thickness is defined as residual film thickness ratio. If it is greater than 95% after processing at 250°C for 1 hour, it is defined to be 20 superior; if it is less than that, it is defined to be inferior.

<Evaluation of transmittance>

After the polyimide pattern is formed, the transmittance between 400 nm to 700 nm is measured. If the transmittance is 25 greater than 90%, it is defined to be superior; if it is less

than that, it is defined to be inferior.

**<Evaluation of chemical resistance>**

Chemical resistance is examined by observing the change  
5 in the thickness when the films are dipped into chemicals (10% aqueous NaOH solution, 10% aqueous HCl solution, and NMP). ((Film thickness before treatment - Film thickness after treatment) / (Film thickness before treatment)) \* 100 (%) is defined to be the ratio of change in thickness. The smaller the 10 ratio of change in thickness when they are treated to chemical substances, the better the chemical resistance. When the ratio of change in thickness after dipping into chemical substances at room temperature for 1 hour is within  $\pm 3\%$ , it is defined to be 15 superior; when it exceeds that range, it is defined to be inferior.

**<Evaluation of planarization>**

In order for films to be used for transparent protection layers for liquid crystal display devices, degree of 20 planarization of the films on topographical substrate is important. Degree of planarization is defined to be (1 - (step height after coating) / (step height before coating)) \* 100 (%). For evaluation of the degree of planarization, 10  $\mu\text{m}$  Line / 10  $\mu\text{m}$  Space (1:1) repetitive pattern with 1.0  $\mu\text{m}$  step height before 25 coating is used. If degree of planarization is greater than 70%,

it is defined to be superior; if it is less than that, it is defined to be inferior.

【Table 1】

Composition	Film thickness (μm)	Heat resistance	Transmittance	Chemical resistance	Degree of planarization	Minimum line width
Preferred Embodiment 1	1.2 μm	Superior	Superior	Superior	Superior	25 μm
Preferred Embodiment 2	1.2 μm	Superior	Superior	Superior	Superior	25 μm
Preferred Embodiment 3	1.2 μm	Superior	Superior	Superior	Superior	30 μm
Comparative Example 1	1.2 μm	Superior	Superior	Inferior	Inferior	50 μm
Comparative Example 2	1.2 μm	Superior	Inferior	Inferior	Inferior	50 μm
Comparative Example 3	1.2 μm	Superior	Superior	Inferior	Superior	40 μm

5

As seen in the above Table 1, transmittance, chemical resistance, and degree of planarization of polyimide films in Preferred Embodiments 1 to 3 of the present invention are superior compared to those of Comparative Examples 1 and 2. It is also 10 seen that the polyimide films show superior properties without additional process compared to Comparative Example 3 which is a conventional method.

#### Industrial Applicability

15

As described in the above, the present invention can provide a negative-type photosensitive polyimide precursor resin

ENDED SHEET(ART. 34)

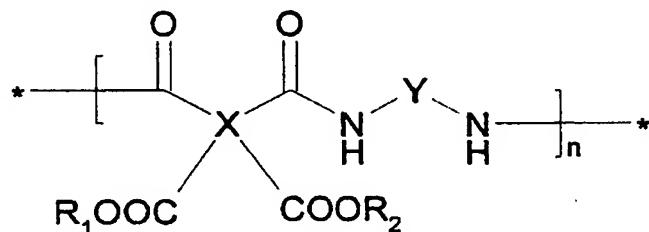
compositions having superior heat resistance, transmittance, chemical resistance, and degree of planarization, and thus is useful in that the above compositions may be applicable for transparent protection layers or insulation layers of liquid crystal display devices.

While the invention has been described in terms of a few preferred embodiments, those skilled in the art will recognize that the invention can be practiced with modification within the spirit and scope of the appended claims.

What is claimed is:

1. A reactive transparent polyimide precursor having the structure represented by the following Chemical Formula 1:

5        <Chemical Formula 1>



wherein X is a tetra-valent organic group derived from alicyclic tetracarboxylic acid dianhydrides having 3 to 30 10 carbon atoms;

Y is a di-valent organic group derived from aliphatic, alicyclic, or non-conjugated aromatic diamines of which number of carbon atoms is 3 to 30 and side chains have one or more ethylenically unsaturated bonds that may be crosslinked by a 15 radical; and

R<sub>1</sub> and R<sub>2</sub> are independently each other hydrogen atom, or organic group having 1 to 20 carbon atoms including one or more ethylenically unsaturated bond(s), provided that they are not hydrogen atom at the same time.

20

2. The reactive transparent polyimide precursor according to Claim 1, wherein the acid value of said reactive transparent

polyimide precursor is within a range of 30 to 200 mg KOH/g.

3. The reactive transparent polyimide precursor according to Claim 1, wherein the molecular weight of said reactive 5 transparent polyimide precursor is within a range of 2,000 to 200,000.

4. A method for preparing the reactive transparent polyimide precursor having the Chemical Formula 1 of Claim 1 10 comprising the steps of:

preparing transparent linear polyamic acid (A) from (a-1) one or more of tetracarboxylic acid dianhydrides selected from alicyclic tetracarboxylic acid dianhydrides having 3 to 30 carbon atoms, and (a-2) one or more of diamines selected from aliphatic, 15 alicyclic, or non-conjugated aromatic diamines of which number of carbon atoms is 3 to 30 and side chains have one or more ethylenically unsaturated bonds that may be crosslinked by a radical; and

esterifying the transparent linear polyamic acid (A) thus 20 obtained with an ethylenically unsaturated compound (B) containing an epoxy group in the same molecule.

5. The method for preparing the reactive transparent polyimide precursors according to Claim 4, wherein the 25 tetracarboxylic acid dianhydride(s) is one or more selected from

the group consisting of 1,2,3,4-cyclobutanetetracarboxylic acid dianhydride (CBDA), 1,2,3,4-cyclopentanetetracarboxylic acid dianhydride (CPDA),

bicyclooct-7-ene-2,3,5,6-tetracarboxylic acid dianhydride

5 (BODA),

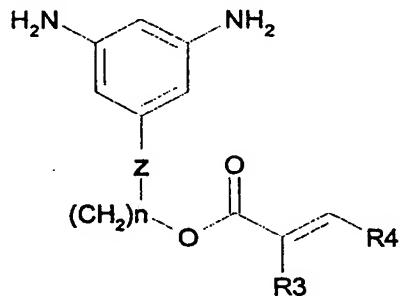
5-(2,5-dioxotetrahydrofuran-3-yl)-3-methylcyclohexene-1,2-dicarboxylic acid anhydride (DOCDA), and 4-(2,5-dioxotetrahydrofuran-3-yl)-tetralin-1,2-dicarboxylic acid anhydride (DOTDA).

10

6. The method for preparing the reactive transparent polyimide precursors according to Claim 4, wherein the diamine(s) is one or more selected from the group consisting of diamines having the general formulae represented by the following

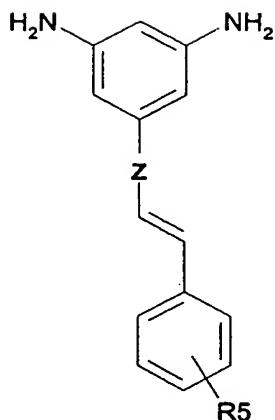
15 Chemical Formulae 7 to 9:

<Chemical Formula 7>

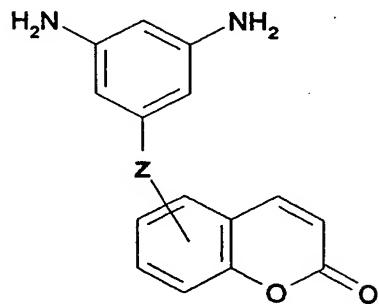


<Chemical Formula 8>

AMENDED SHEET(ART. 34)



<Chemical Formula 9>



5 wherein Z is one of ester, amide, imide, ether, and carbonyl group;

R<sub>3</sub>, R<sub>4</sub>, and R<sub>5</sub> are independently each other hydrogen atom or alkyl or aryl group having 1 to 20 carbon atoms; and n is an integer of 1 to 20.

10

7. The method for preparing the reactive transparent polyimide precursors according to Claim 4, wherein the diamine(s) is one or more selected from the group consisting of 2-(methacryloyloxy)ethyl 3,5-diaminobenzoate,

3,5-diaminophenyl cinnamate, and coumaronyl  
3,5-diaminobenzoate.

8. The method for preparing the reactive transparent  
5 polyimide precursors according to Claim 4, wherein the  
ethylenically unsaturated compound (B) containing an epoxy group  
in the same molecule is one or more compounds selected from the  
group consisting of allyl glycidyl ether, glycidyl acrylate,  
glycidyl methacrylate, 3,4-epoxycyclohexylmethyl acrylate,  
10 3,4-epoxycyclohexylmethyl methacrylate, glycidyl  
5-norbornene-2-carboxylate (a mixture of endo and exo forms),  
glycidyl 5-norbornene-2-methyl-2-carboxylate (a mixture of  
endo and exo forms), 1,2-epoxy-5-hexene, and  
1,2-epoxy-9-decene.

15

9. A photosensitive polyimide precursor resin composition,  
characterized in that it is prepared by using the reactive  
transparent polyimide precursor of Claim 1 and one or more  
photo-initiators as essential components, and adding one or more  
20 compounds selected from the group consisting of photosensitizers,  
multi-functional monomers, and common coating additives if  
necessary.

10. The photosensitive polyimide precursor resin  
25 composition according to Claim 9, wherein the weight of the

reactive transparent polyimide precursor is in the range of 10 to 99 weight % relative to the weight of the total solids and the weight of the photo-initiators is in the range of 0.1 to 90 weight % relative to the weight of the total solids.

5

11. The photosensitive polyimide precursor resin composition according to Claim 9, wherein the thickness of polyimide film obtained from the photosensitive polyimide precursor resin composition is in the range of 0.5 to 100  $\mu\text{m}$ .

10

12. The photosensitive polyimide precursor resin composition according to Claim 9, wherein the thermal decomposition temperature of polyimide film obtained from the photosensitive polyimide precursor resin composition is within 15 a range of 300 to 500°C.

13. The photosensitive polyimide precursor resin composition according to Claim 9, wherein the transmittance between 400 to 700 nm of polyimide film obtained from the 20 photosensitive polyimide precursor resin composition is 90% or higher.

14. The photosensitive polyimide precursor resin composition according to Claim 9, wherein the dielectric constant 25 of polyimide film measured at 1 kHz, obtained from the

photosensitive polyimide precursor resin composition, is within a range of 2.5 to 4.0.

15. A photosensitive transparent protection layer or  
5 insulation layer, characterized in that it is prepared by using  
the photosensitive polyimide precursor resin composition  
according to Claim 9.

16. A liquid crystal display device, characterized in that  
10 the photosensitive polyimide precursor resin composition  
according to Claim 9 is applied to transparent protection  
layer(s) or insulation layer(s).

17. A liquid crystal display device, characterized in that  
15 the photosensitive polyimide precursor resin composition  
according to Claim 9 is applied as organic insulation materials  
for liquid crystal display device.

**Abstract**

The present invention relates to an aqueous alkali-developable photosensitive polyimide precursor resin 5 composition that is appropriate for highly heat-resistant transparent protection layers and insulation layers for liquid crystal display devices.

In more detail, the present invention relates to a reactive transparent polyimide precursor represented in the following 10 Chemical Formula 1, wherein it is prepared by following steps of preparing transparent linear polyamic acid (A) from (a-1) one or more of tetracarboxylic acid dianhydrides selected from alicyclic tetracarboxylic acid dianhydrides having 3 to 30 carbon atoms, and (a-2) one or more of diamines selected from aliphatic, 15 alicyclic, or non-conjugated aromatic diamines of which number of carbon atoms is 3 to 30 and side chains have one or more ethylenically unsaturated bonds that may be crosslinked by a radical; and esterifing the transparent linear polyamic acid (A) thus obtained with an ethylenically unsaturated compound 20 (B) containing an epoxy group in the same molecule, and an aqueous alkali developable negative-type photosensitive transparent polyimide precursor resin composition prepared by using the same as a main component. The photosensitive transparent polyimide precursor resin compositions according to the present invention 25 have a superior photosensitivity, and thus, may be used for

transparent protection layers and insulation layers of liquid crystal display devices having superior heat resistance, chemical resistance, mechanical strength, and electricity insulation.

5

<Chemical Formula 1>

